

**Amendments to the Claims:**

This listing of claims will replace all prior versions, and listings, of claims in the captioned application:

**Listing of Claims:**

Claim 1-17 (cancelled).

Claim 18 (previously presented): A method for enhancing the absorption rate of a pharmaceutically acceptable amine into the blood of a human comprising administering a stable suspension comprising (i) a pharmacologically effective dosage of a pharmaceutically acceptable amine and (ii) an effective amount of ibuprofen.

Claim 19 (previously presented): The method of claim 18 wherein the amine is pseudoephedrine.

Claim 20 (previously presented): The method of claim 18 wherein the enhanced absorption is indicated by AUC 1 H (early drug exposure) that is at least about 10% higher than the early drug exposure of the same amine from a single-ingredient liquid.

Claim 21 (previously presented): The method of claim 18 wherein the enhanced absorption is indicated by AUC 2 H (early drug exposure) that is at least about 10% higher than the early drug exposure of the same amine from a single-ingredient liquid.

Claim 22 (previously presented): The method of claim 18 wherein the enhanced absorption is indicated by a C MAX (maximum or peak concentration) that is at least about 10% greater than the C MAX of the same amine from a single-ingredient liquid.

Claim 23 (previously presented): The method of claim 18 wherein the amine is provided in a range from about 15 mg to about 60 mg per dosage unit.

Claim 24 (previously presented): The method of claim 18 wherein the ibuprofen is provided in an amount of from about 40 mg to about 800 mg per dosage unit.

Claim 25 (previously presented): The method of claim 23 wherein the ibuprofen is provided in an amount of from about 40 mg to about 800 mg per dosage unit.

Claim 26 (previously presented): The method of claim 18 wherein the amine is provided at about 15 milligrams per 5 mL and the ibuprofen is provided at about 100 milligrams per 5 mL.

Claim 27 (previously presented): The method of claim 18 wherein the human is a child.

Claims 28-41 (cancelled).

Claim 42 (currently amended): A stable suspension comprising (i) a pharmacologically effective amount of a pharmaceutically acceptable amine and (ii) a pharmacologically effective amount of ibuprofen. ~~The suspension of claim 30,~~ wherein the suspension further comprises xanthan gum, pregelatinized starch, polyoxyethylene sorbitan monooleate and a taste masking agent selected from the group consisting of sugar, sweet polyhydric alcohol, cyclamates, aspartame, sucralose saccharin, flavoring agents and mixtures thereof.